Claims:

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- 1. A pharmaceutical composition for use in decreasing DNA damage comprising an effective daily dose of about 0.1 to 20 mg lutein, and at least one of the group consisting of beta-carotene and lycopene in amounts sufficient to act synergistically with lutein.
- 2. The pharmaceutical composition of claim 1, wherein the composition further comprises at least one of about 0.1 mg to 20 mg beta-carotene or about 0.1 to 20 mg lycopene.
 - 3. The pharmaceutical composition of claim 1, wherein the composition further comprises a lipophilic component.
 - 4. The pharmaceutical composition of claim 1, wherein the composition further comprises a carotenoid-containing dry powder in the form of a multicore structure in which at least two cores of a multicore structure comprise one or more different carotenoids of the group consisting of substantially purified lutein, beta-carotene, and lycopene.
 - 5. The pharmaceutical composition of claim 4, wherein the carotenoid-containing dry powder is formed into at least one of drink preparations, tablets, sugar coated tablets, hard gelatin capsules, soft gelatin capsules, and cellulose capsules.
 - 6. A nutritional composition suitable for use in protecting against a free radical associated disorder, comprising a daily dose of at least two carotenoids selected from the group consisting of substantially purified lutein, beta-carotene, and lycopene.
 - 7. The nutritional composition of claim 6, wherein the daily dose of at least two carotenoids is selected from about 0.1% to 50% by weight beta-carotene, about 0.1% to 50% by weight lycopene, and about 0.1% to 50% by weight lutein.

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- 8. A method of decreasing oxidative damage in a subject comprising: administering a synergistic combination of carotenoids to the subject, wherein the synergistic combination comprises at least two carotenoids selected from the group consisting of lutein, beta-carotene, and lycopene.
- 9. The method of claim 8, wherein the synergistic combination of carotenoids is selected from the group consisting of a daily unit dose of about 0.1 mg to 20 mg beta-carotene, about 0.1 to 20 mg lycopene, and about 0.1 to 20 mg lutein to the subject.
- 10. The method of claim 8, wherein the method comprises administering about 0.5 mg to 10 mg beta-carotene, about 0.5 to 10 mg lycopene, and about 0.5 to 10 mg lutein to the subject.
- 11. The method of claim 8, wherein the synergistic combination of carotenoids is selected from the group consisting of a daily unit dose of about 1 part of betacarotene, 0.02 to 20 parts of lycopene and 0.02 to 20 parts of lutein.
- 20 12. The method of claim 8, wherein the synergistic combination of carotenoids is selected from the group consisting of a daily unit dose of about 1 part of beta-carotene, 0.1 to 2 parts of lycopene and 0.1 to 2 parts of lutein.
- 13. The method of claim 8, wherein the method further comprises administering a lipophilic component, such that antioxidant capacity in the aqueous and lipid compartments of plasma is increased.
 - 14. The method of claim 8, wherein the method further comprises administering a carotenoid-containing dry powder in the form of a multicore structure in which at least two cores of a multicore structure comprise one or more different carotenoids of the group consisting of substantially purified lutein, beta-carotene, and lycopene.

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- 15. The method of claim 14, wherein the method comprises administering the carotenoid-containing dry powder in a form selected from the group consisting of drink preparations, tablets, sugar coated tablets, hard gelatin capsules, soft gelatin capsules, and cellulose capsules.
- 16. A method of reducing effects of aging in a subject comprising: administering a synergistic combination of carotenoids to the subject, wherein the synergistic combination comprises at least two of the group consisting of lutein, beta-carotene, and lycopene, whereby DNA damage in the subject is decreased thereby reducing the effects of aging.
- 17. The method of claim 16, wherein the synergistic combination of carotenoids is selected from the group consisting of a daily unit dose of about 0.1 mg to 20 mg beta-carotene, about 0.1 to 20 mg lycopene, and about 0.1 to 20 mg lutein to the subject.
 - 18. The method of claim 16, wherein the method comprises administering about 0.5 mg to 10 mg beta-carotene, about 0.5 to 10 mg lycopene, and about 0.5 to 10 mg lutein to the subject.
 - 19. The method of claim 16, wherein the method comprises a carotenoid-containing dry powder in the form of a multicore structure in which at least two cores of a multicore structure comprise one or more different carotenoids of the group consisting of substantially purified lutein, beta-carotene, and lycopene.
 - 20. The method of claim 19, wherein the method comprises administering the carotenoid-containing dry powder in a form selected from the group consisting of drink preparations, tablets, sugar coated tablets, hard gelatin capsules, soft gelatin capsules, and cellulose capsules.